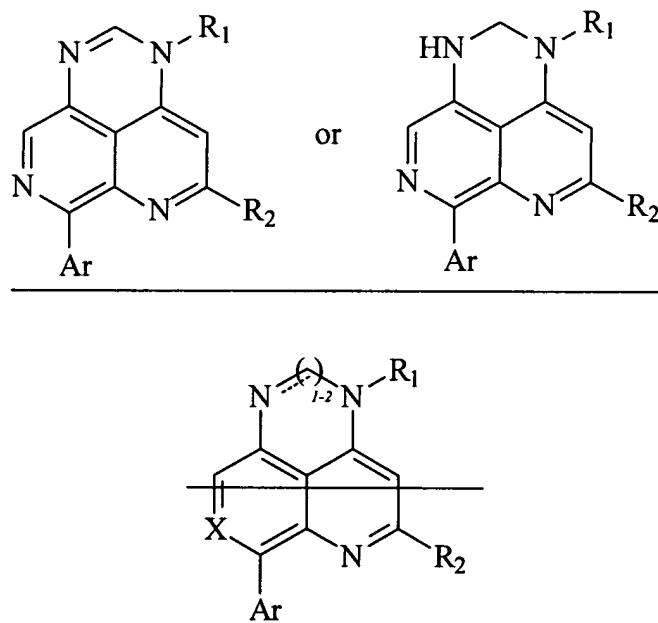


Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

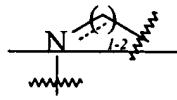
Listing of Claims:

1. (Amended) A compound having the following structure:



including stereoisomers and pharmaceutically acceptable salts thereof,

wherein:



~~represents $\text{N}=\text{CH}-$, $\text{NH}-\text{CH}_2-$ or $\text{NH}-(\text{CH}_2)_2-$~~

~~X is N or CR_3~~ ;

R₁ is $-\text{CH}(\text{R}_4)(\text{R}_5)$;

R₂ is C₁₋₆alkyl;

R₃ is hydrogen or C₁₋₆alkyl;

R₄ is hydrogen, C₁₋₆alkyl, mono- or di(C₃₋₆cycloalkyl)methyl, C₃₋₆cycloalkyl, C₃₋₆alkenyl, hydroxyC₁₋₆alkyl, C₁₋₆alkylcarbonyloxyC₁₋₆alkyl, or C₁₋₆alkyloxyC₁₋₆alkyl, and

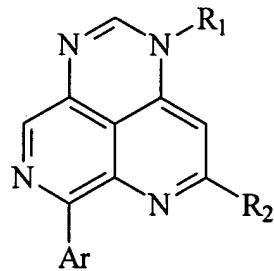
R₅ is C₁₋₈alkyl, mono- or di(C₃₋₆cycloalkyl)methyl, Ar¹CH₂, C₃₋₆alkenyl, C₁₋₆alkyloxyC₁₋₆alkyl, hydroxyC₁₋₆alkyl, thienylmethyl, furanylmethyl, C₁₋₆alkylthioC₁₋₆alkyl, morpholinyl, mono- or di(C₁₋₆alkyl)aminoC₁₋₆alkyl, di(C₁₋₆alkyl)amino, C₁₋₆alkylcarbonylC₁₋₆alkyl, C₁₋₆alkyl substituted with imidazolyl, -CH₂Obenzyl, or a radical of the formula -(C₁₋₆alkanediyl)-O-CO-Ar¹,

or R₄ and R₅ taken together with the carbon atom to which they are bonded form a C₅₋₈cycloalkyl optionally substituted with one or more substituents independently selected from C₁₋₆alkyl;

Ar is phenyl substituted with 1, 2 or 3 substituents independently selected from halo, C₁₋₆alkyl, trifluoromethyl, cyano, C₁₋₆alkyloxy, benzyloxy, C₁₋₆alkylthio, nitro, amino, and mono- or di(C₁₋₆alkyl)amino; or an aromatic C₃₋₁₂heterocycle optionally substituted with 1, 2 or 3 substituents independently selected from halo, C₁₋₆alkyl, trifluoromethyl, hydroxy, cyano, C₁₋₆alkyloxy, benzyloxy, C₁₋₆alkylthio, nitro, amino, mono- or di(C₁₋₆alkyl)amino, and piperidinyl; and

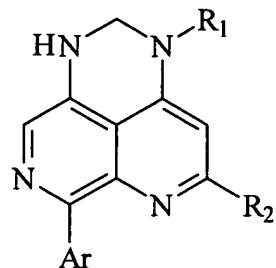
Ar¹ is phenyl, pyridinyl, or phenyl substituted with 1, 2 or 3 substituents independently selected from halo, C₁₋₆alkyl, C₁₋₆alkyloxy, di(C₁₋₆alkyl)aminoC₁₋₆alkyl, trifluoromethyl and C₁₋₆alkyl substituted with morpholinyl.

2. (Amended) The compound of claim 1 having the structure:



3. (Canceled)

4. (Amended) The compound of claim 1 having the structure:



- 5.-7. (Canceled)
8. (Original) The compound of claim 1 wherein Ar is 2,4-dichlorophenyl.
9. (Original) The compound of claim 1 wherein Ar is 2-chloro-4-methylphenyl.
10. (Original) The compound of claim 1 wherein Ar is 2-methyl-4-chlorophenyl.
11. (Original) The compound of claim 1 wherein Ar is 2,4,6-trimethylphenyl.
12. (Original) The compound of claim 1 wherein Ar is 2-chloro-4-methoxyphenyl.
13. (Original) The compound of claim 1 wherein Ar is 2-methyl-4-methoxyphenyl.

14. (Original) The compound of claim 1 wherein Ar is 2,4-dimethoxy-phenyl.
15. (Original) The compound of claim 1 wherein Ar is 4-dimethylamino-2-methyl-3-pyridyl.
16. (Original) The compound of claim 1 wherein Ar is 4-dimethylamino-6-methyl-3-pyridyl.
17. (Original) The compound of claim 1 wherein Ar is 4-dimethylamino-3-pyridyl.
18. (Original) The compound of claim 1 wherein R₁ is -CH(n-propyl)₂.
19. (Original) The compound of claim 1 wherein R₁ is -CH(n-propyl)(CH₂OCH₃).
20. (Original) The compound of claim 1 wherein R₁ is -CH(benzyl)(CH₂OCH₃).
21. (Original) The compound of claim 1 wherein R₁ is -CH(CH₂OR)₂ and each occurrence of R is independently selected from C₁₋₆alkyl.
22. (Original) The compound of claim 1 wherein R₁ is -CH(CH₂OR)(ethyl) and each occurrence of R is independently selected from C₁₋₆alkyl.
23. (Original) The compound of claim 1 wherein R₁ is -CH(CH₂OR)(n-butyl) and each occurrence of R is independently selected from C₁₋₆alkyl.

24. (Original) The compound of claim 1 wherein R₁ is -CH(CH₂OR)(tert-butyl) and each occurrence of R is independently selected from C₁₋₆alkyl.

25. (Original) The compound of claim 1 wherein R₁ is -CH(CH₂OR)(4-chloro-benzyl) and each occurrence of R is independently selected from C₁₋₆alkyl.

26. (Original) The compound of claim 1 wherein R₁ is -CH(CH₂OR)(CH₂CH₂SCH₃) and each occurrence of R is independently selected from C₁₋₆alkyl.

27. (Original) The compound of claim 1 wherein R₁ is -CH(CH₂CH₃)(CH₂Obenzyl).

28. (Original) The compound of claim 1 wherein R₂ is methyl.

29. (Original) The compound of claim 1 wherein R₂ is ethyl.

30. (Amended) A pharmaceutical composition comprising a compound of claim 1 in combination with a pharmaceutically acceptable carrier or diluent.

31. (Amended) A method for treating stroke, anxiety, depression or irritable bowel syndrome ~~a disorder manifesting hypersecretion of CRF~~ in a warm-blooded animal, comprising administering to the animal an effective amount of the pharmaceutical composition of claim 30.

32. (Original) The method of claim 31 wherein the disorder is stroke.

33. (Original) The method of claim 31 wherein the disorder is anxiety.

34. (Original) The method of claim 31 wherein the disorder is depression.

35. (Original) The method of claim 31 wherein the disorder is irritable bowel syndrome.